HIV/AIDS Epidemiology Program

HEALTH

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Concurrent Diagnoses of HIV and AIDS in New Mexico

Introduction

The HIV/AIDS Epidemiology Program in New Mexico conducts confidential name-based **HIV/AIDS** reporting and subsequent epidemiological analyses. The HIV surveillance case definition includes a positive screening test result (e.g., reactive EIA) confirmed by a positive result from a supplemental HIV antibody test (e.g., Western Blot). HIV infected persons who have progressed to AIDS are identified based on a case definition using either immunological data (specifically, an absolute CD4⁺ T lymphocyte cell count of <200 or CD4⁺ <14% of total lymphocytes) and/or a clinical diagnosis of an opportunistic infection (OI).^{1,2}

With the advent of highly active antiretroviral therapy (HAART) in 1996 and proper preventive therapy for Ols, the progression of HIV to AIDS can be prevented or delayed considerably. allowing persons infected with HIV to live longer, fuller lives. In this report, we use New Mexico HIV/AIDS surveillance data collected from 1998 to 2008 to make inferences concurrently diagnosed regarding cases. defined as those cases that progress to AIDS less than one year after their initial HIV Concurrently diagnosis. diagnosed represent persons who are diagnosed late in the course of infection, i.e. 'late testers', and indicate missed opportunities to improve the health outcomes of HIV+ persons.

Concurrent diagnoses in New Mexico

Despite our efforts to expand and increase HIV testing and provide patient education, people continue to be diagnosed late in the course of their infection. Table 1 presents HIV/AIDS data in New Mexico from 1998 to 2008 for various population groups. Among New Mexicans diagnosed with HIV, 43.2% (646/1,497) received a concurrent AIDS diagnosis; this percentage is relatively higher than the U.S. concurrent diagnosis of approximately 38%.³

The spectrum of 'late testers' identified in Table 1 is clearly as diverse as the entire population reported with HIV/AIDS. In New Mexico, the proportion of HIV positive Hispanics with concurrent diagnoses (46%) was higher than that of Whites, African Americans, and American Indian/Alaska Natives (the number of Asians and those identifying as multi-race were relatively small for accurate comparison). The higher proportion found among Hispanics is concordant with the CDC's findings through case surveillance in 30 U.S. states as well as Supplement through the to HIV/AIDS Surveillance (SHAS) interviews conducted in New Mexico and across the U.S. in 2000-2003.4

proportion of persons concurrently diagnosed increased across age categories. For example, more than half (53.8%) of persons aged 40-49 at the time of HIV diagnosis were concurrently diagnosed with AIDS: proportion was 63.6% for persons aged 50-59 and 73.1% for persons aged >60 years. A smaller proportion of women received concurrent diagnoses relative to men (34.4% vs. 44.6%, respectively). Concurrent diagnoses were highest among residents of the southeast and northeast regions of the state (52.1% and 48.3%, respectively; Figure 1).

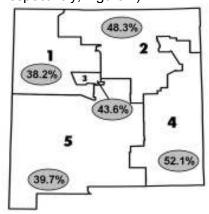


Figure 1. New Mexico DOH Public Health Regions and associated concurrent diagnoses percentages, 1998 - 2008

Figure 2 depicts the annual incidence of concurrent and non-concurrent diagnoses among persons with HIV/AIDS in New Mexico since 1998. This figure illustrates that, though the number of concurrently diagnosed cases has remained relatively constant, progress has been made in reducing the proportion of persons diagnosed with HIV who are concurrently diagnosed with AIDS. Between 1998 and 2004, there were 45.6% concurrent cases among new diagnoses. Between 2005 and 2008, this average declined to 39.5%.

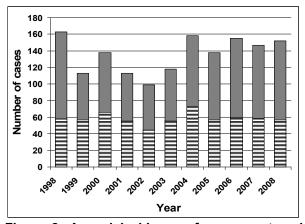


Figure 2. Annual incidence of concurrent and non-concurrent diagnoses among persons with HIV/AIDS in New Mexico. The hatched areas represent concurrent diagnoses, whereas the solid areas represent non-concurrent HIV/AIDS diagnoses.

Concurrent diagnoses and morbidity

Among reported HIV cases that progressed to AIDS, the AIDS-defining condition among those concurrently diagnosed is more often based on the presence of immunologic results than on clinical features. Figure 3 depicts New Mexico AIDS cases that concurrently and non-concurrently diagnosed by immunological or clinical AIDS diagnosis. Clearly, most concurrent cases had their AIDS-defining condition identified via reported CD4+ levels (filled circles). Also, it is evident that, since 2003, there has been an increased disparity between concurrent and non-concurrent cases (closed circles vs. open circles and open squares) as well as between the AIDS-defining condition; these trends could represent a bias toward under-reporting of OI data among those non-concurrently diagnosed coupled with relatively improved reporting of laboratory (immunological) data.

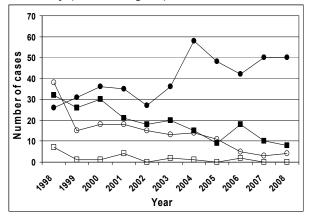


Figure 3. From 1998-2008, concurrently and nonconcurrently diagnosed AIDS immunological and clinical criteria. Circles immunological represent (CD4+) **AIDS** diagnoses, whereas squares represent AIDS cases identified by opportunistic infections. Within each symbol type, the open and closed represent non-concurrent concurrent diagnoses, respectively.

Among all cases that have developed an OI, those cases concurrently diagnosed developed an OI relatively sooner than those cases non-concurrently diagnosed (Figure 4). Specifically, greater than 90% of concurrent cases developed an OI within 36 months of diagnosis, including at the time of diagnosis. In contrast, less than 25% of non-concurrent cases developed an OI within 36 months.

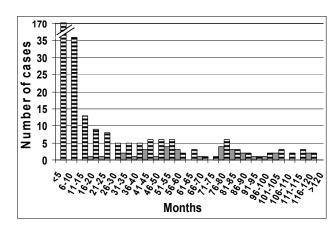


Figure 4. Months from AIDS diagnosis to acquire an OI. The hatched areas represent concurrent diagnoses, whereas the solid areas represent non-concurrent AIDS diagnoses.

In examining morbidity associated with concurrently vs. non-concurrently diagnosed cases, it's important to consider which OIs affect persons with HIV. Opportunistic infections are caused by many types of pathogens, including protozoa, bacteria, fungi and viruses.⁵ Without effective prophylaxis, infection with compromises the immune system, allowing OIs to develop and cause health problems, potentially causing premature mortality. Of New Mexico AIDS cases that ever developed an OI, 89.7% (269/300) were concurrently diagnosed, whereas only 10.3% (31/300) were not concurrently diagnosed. Table 2 lists the frequency (by concurrent vs. non-concurrent status) of the most common Ols.

	Concurrent	Non- concurrent	
Opportunistic Infection	n	n	
Kaposi's sarcoma	29	3	
Pneumocystosis	111	11	
Wasting syndrome	30	1	
Toxoplasmosis	5	0	
Candidiasis	38	5	
Mycobacterium avium	11	3	
Mycobacterium tuberculosis	16	1	
Cytomegalovirus	26	5	
Coccidioidomycosis	7	1	
Cryptosporidiosis	5	0	
Herpes simplex	7	1	
Histoplasmosis	8	0	
HIV encephalopathy	7	0	
Lymphoma	8	0	
Pneumonia	8	0	
Other	26	7	

Table 2. Counts of common opportunistic infections among New Mexico AIDS cases concurrently and non-concurrently diagnosed, 1998-2008. Some cases had more than one OI.

Public Health Implications

The public health implications of concurrently-diagnosed individuals, or 'late testers', are three-fold. First, persons diagnosed with HIV when they are already symptomatic and/or immunosuppressed may miss opportunities to delay the progression of HIV/AIDS disease by timely initiation of anti-retroviral treatment. Second, given that late diagnosis may be an

important contributing factor to morbidity, concurrent diagnoses hinder the administration of prophylaxis for opportunistic infections. Third, because they are unaware of their HIV status, 'late testers' may unknowingly transmit HIV to others.

By identifying New Mexico populations disparately affected by concurrent diagnoses, we may gain further insight into testing behaviors among specific groups. Our New Mexico-specific findings suggest the need for targeted early testing for: 1) males; 2) Hispanics; and 3) at-risk residents of the southeast and northeast regions of the state. Ultimately, these data can be used to refine our HIV testing and prevention strategies to better serve these identified at-risk populations.

References

- **1.** Kaplan, JE (2000) Epidemiology of human immunodeficiency virus-associated opportunistic infections in the United States in an era of highly active antiretroviral therapy. *Clinical Infectious Disease*; 30:S5-14
- **2.** Kaplan, JE, H Masur, KK Holmes (2002) Guidelines for preventing opportunistic infections among HIV-infected presons 2002. Recommendations of the U.S. Public Health Service and the Infectious Diseases Society of America. *MMWR Recomm.* Rep 51:1-52
- **3.** CDC (2009) Late HIV testing 34 states, 1996-2005. *MMWR Morb Mortal Wkly Rep* Jun 26; 58(24): 661-5
- **4.** CDC (2009) Cases of HIV Infection and AIDS in the United States and Dependent Areas, by Race/Ethnicity, 2002–2006. HIV/AIDS Surveillance Supplemental Report 13(1) http://www.cdc.gov/hiv/topics/surveillance/resources/reports/2008supp vol13no1/default.htm
- **5.** NMDOH, HIV & Hepatitis Epidemiology Program (2009) HIV/AIDS and Opportunistic Infections. Winter Quarterly Report, January.

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TABLE 1. Concurrent Diagnoses, by selected characteristics, NM (1998-2008) vs. US[†] (1998-2005)

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	No. with HIV/AIDS diagnosis in NM		NM AIDS diagnosis ≤ 1 year after HIV diagnosis		No. with HIV diagnosis in US		US AIDS diagnosis ≤ 1 year after HIV diagnosis	
	N	% of total	N	% within group	N	% of total	% within group	
Sex								
Male	1,273	85.0%	569	44.6%	200,882	71.4%	40.2%	
Female	224	15.0%	77	34.4%	80,539	28.6%	34.8%	
Race / Ethnicity								
White	493	32.9%	206	41.8%	85,149	30.3%	37.1%	
Hispanic	758	51.5%	349	46.0%	42,098	15.0%	42.0%	
American Indian / Alaskan Native	136	8.9%	52	38.2%	1,463	0.5%	39.0%	
African American	90	5.7%	30	33.0%	148,146	52.6%	38.7%	
Asian/Pacific Islander	10	0.7%	5	50.0%	1,742	0.6%	44.1%	
Multirace	10	0.7%	4	40.0%				
Age at Diagnosis								
< 13	4	0.3%	2	50.0%	2,100	0.7%	19.0%	
13-19	44	2.9%	2	4.5%	8,303	3.0%	16.1%	
20-29	354	23.6%	72	20.3%	60,089	21.4%	24.7%	
30-39	497	33.2%	223	44.9%	99,657	35.4%	38.5%	
40-49	392	26.2%	211	53.8%	74,934	26.6%	45.3%	
50-59	154	10.3%	98	63.6%	26,989	9.6%	51.3%	
60+	52	3.5%	38	73.1%	9,349	3.3%	57.0%	
Danian of Diamonia								
Region at Diagnosis	044	40.40/	00	20.20/	NI/A	NI/A	NI/A	
1, Northwest	241	16.1%	92	38.2%	N/A	N/A	N/A	
2, Northeast	205	13.7%	99	48.3%	N/A	N/A	N/A	
3, Bernalillo County	675	45.1%	294	43.6%	N/A	N/A	N/A	
4, Southeast	94	6.3%	49	52.1%	N/A	N/A	N/A	
5, Southwest	282	18.8%	112	39.7%	N/A	N/A	N/A	
Mode of Exposure				-				
Males - Sex with Males (MSM)	776	51.8%	341	43.9%	100,231	35.6%	40.9%	
Injection Drug User (IDU) - Males	105	7.0%	47	44.8%	20,970	7.5%	42.0%	
Injection Drug User (IDU) - Females	47	3.1%	18	38.3%	11,184	4.0%	32.9%	
MSM/IDU	107	7.1%	46	43.0%	9,494	3.4%	38.2%	
Heterosexual - Males	71	4.7%	36	50.7%	21,883	7.8%	43.1%	
Heterosexual - Females	111	7.4%	41	36.9%	39,218	13.9%	34.0%	
Other or Unknown	280	18.7%	117	41.8%	78,441	27.9%	37.7%	
Year of diagnosis								
1998	163	10.9%	58	35.6%	15,429	5.5%	41.6%	
1999	113	7.5%	57	50.4%	23,295	8.3%	40.6%	
2000	138	9.2%	66	47.8%	28,842	10.2%	39.5%	
2001	113	7.5%	56	49.6%	38,818	13.8%	36.4%	
2002	99	6.6%	45	45.5%	36,244	12.9%	36.7%	
2003	118	7.9%	56	47.5%	33,826	12.0%	37.7%	
2004	158	10.6%	73	46.2%	35,645	12.7%	37.6%	
2005	138	9.2%	57	41.3%	34,424	12.7%	36.4%	
2006	155	10.4%	60	38.7%	**	**	**	
2007	147	9.8%	60	40.8%	**	**	**	
2008	155	10.4%	58	37.4%	**	**	**	
2000	133	10.4 /0	50	37.470				